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## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## Listing of Claims:

Claims 1-18 (Canceled)

- 19. (Previously Amended) A method of determining the presence of one or more target analytes in one or more samples comprising:
- a) adding said one or more samples to a first substrate comprising a plurality of assay locations, such that said one or more samples is contained at a plurality of said assay locations;
  - b) contacting said one or more samples with a second substrate comprising:
- i) a plurality of array locations, each array location comprising a plurality of discrete sites, wherein at least one assay location is in fluid contact with at least one array location; and
- ii) a population of microspheres comprising at least a first and a second subpopulation each comprising a bioactive agent, wherein said microspheres are distributed on at least one of said array locations such that said discrete sites each contain no more than one microsphere; and
  - c) determining the presence or absence of said target analyte.
- 20. (Currently Amended) A method according to claim [[18]] 19 or 36, wherein each of said assay locations comprises a library of bioactive agents.
- 21. (Canceled)
- 22. (Currently Amended) A method according to claim [[18]] 19 or 36, wherein each discrete site is a bead well.
- 23-24. (Canceled)
- 25. (Currently Amended) A method according to claim 19 or 36, wherein said first substrate is a microtiter plate.

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- 26. (Currently Amended) A method according to claim 19 or [[25]] <u>36</u>, wherein said second substrate comprises a plurality of fiber optic bundles comprising a plurality of individual fibers, each bundle comprising an array location, and each individual fiber comprising a bead well.
- 27. (Currently Amended) A method according to claim 19 or 36, wherein each of said subpopulations further comprise an optical signature capable of identifying said bioactive agent.
- 28. (Currently Amended) A method according to claim 19 or 36, wherein each of said subpopulations further comprise an identifier binding ligand that will bind a decoder binding ligand such that the identification of the bioactive agent can be elucidated.
- 29. (Currently Amended) A method according to claim 18 or 19 or 36, wherein at least one of said target analytes is a nucleic acid.
- 30. (Currently Amended) A method according to claim 18 or 19 or 36, wherein said microspheres are randomly distributed on said surface.
- 31. (Currently Amended) A method according to claim 18 or 19 or 36, wherein at least a first subpopulation of microspheres comprises a bioactive agent comprising nucleic acids.
- 32. (Currently Amended) A method according to claim 18 or 19 or 36, wherein at least a first subpopulation of microspheres comprises a bioactive agent comprising a protein.
- 33. (Previously presented) A method according to claim 20, wherein at least a first and second of said assay locations comprise the same library of bioactive agents.
- 34. (Previously presented) A method according to claim 20, wherein at least a first and second of said assay locations comprise different libraries of bioactive agents.
- 35. (Canceled)
- 36. (Previously presented) A method of determining the presence of one or more target analytes in one or more samples comprising:
- a) adding said one or more samples to a first substrate comprising a plurality of assay locations, such that said one or more samples is contained at a plurality of said assay locations;

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b) contacting said one or more samples with a second substrate comprising:

- i) a composite array comprising a plurality of array locations, each array location comprising a plurality of discrete sites, wherein at least one assay location is in fluid contact with at least one array location; and
- ii) a population of microspheres comprising at least a first and a second subpopulation each comprising a bioactive agent, wherein said microspheres are distributed on said surface such that said discrete sites each contain no more than one microsphere; and
  - c) determining the presence or absence of said target analyte.